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B'

R<sub>3</sub> is H, -SO<sub>2</sub> (C<sub>1-6</sub> alkyl), -SO<sub>2</sub> phenyl, (C=O)(C<sub>1-6</sub> alkyl), or -W'Z';

W' is a covalent bond, (C=O), SO<sub>2</sub>, or C<sub>1-6</sub> alkyl;

Z' is C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, C<sub>3-8</sub> cycloalkyl, ~~phenyl~~, or a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrol, imidazol, pyrazol, isothiazol, isoxazol, pyrid, pyrazin, pyrimidin, pyridazin, indolizin, isoindol, indol, indazol, purin, quinol, furazan, pyrrolidin, pyrrolin, imidazolidin, imidazolin, pyrazolidin, pyrazolin, piperid, piperazin, indolin, and morpholinC<sub>2-6</sub> ~~heterocyclic radical, optionally including in the ring up to 3 additional heteroatoms or moieties independently selected from O, N, NH, S, SO, and SO<sub>2</sub>~~; or Z' is NR<sub>13</sub>R<sub>14</sub> where each of R<sub>13</sub> and R<sub>14</sub> is independently selected from C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl, phenyl, benzyl, C<sub>3-8</sub> cycloalkyl, and a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrol, imidazol, pyrazol, isothiazol, isoxazol, pyrid, pyrazin, pyrimidin, pyridazin, indolizin, isoindol, indol, indazol, purin, quinol, furazan, pyrrolidin, pyrrolin, imidazolidin, imidazolin, pyrazolidin, pyrazolin, piperid, piperazin, indolin, and morpholinC<sub>2-5</sub> ~~heterocyclic radical~~; each of R<sub>5</sub>, R<sub>6</sub>, R<sub>7</sub> and R<sub>8</sub> is independently H, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, halo, nitro, or amino;

one of R<sub>a</sub>, R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub>, and R<sub>e</sub> is WZ and the others are independently selected from H, C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, halo, nitro, and amino;

W is -O-, R<sub>9</sub>, O-R<sub>9</sub>, NR<sub>10</sub>, -(CO)(O)R<sub>9</sub>, -O (CO)R<sub>9</sub>,

-(CO)NR<sub>10</sub>, or -N(R<sub>10</sub>)-CO-R<sub>9</sub>, wherein R<sub>9</sub> is C<sub>1-6</sub> alkylene, C<sub>2-6</sub> alkynylene, C<sub>2-6</sub> alkenylene, phenylene, or C<sub>2-5</sub> a heterocyclic bivalent radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrol, imidazol, pyrazol, isothiazol, isoxazol, pyrid, pyrazin, pyrimidin, pyridazin, indolizin, isoindol, indol, indazol, purin, quinol, furazan, pyrrolidin, pyrrolin, imidazolidin, imidazolin, pyrazolidin, pyrazolin, piperid, piperazin, indolin, and morpholin, and R<sub>10</sub>

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is H, C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkynyl, C<sub>2-6</sub> alkenyl, phenyl, or a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl, isoxazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indoliziny, isoindolyl, indolyl, indazolyl, purinyl, quinolyl, furazanyl, pyrrolidinyl, pyrrolinyl, imidazolidinyl, imidazoliny, pyrazolidinyl, pyrazolinyl, piperidyl, piperazinyl, indolinyl, and morpholinyl C<sub>2-5</sub> heterocyclic radical;

Z is a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl, isoxazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indoliziny, isoindolyl, indolyl, indazolyl, purinyl, quinolyl, furazanyl, pyrrolidinyl, pyrrolinyl, imidazolidinyl, imidazoliny, pyrazolidinyl, pyrazolinyl, piperidyl, piperazinyl, indolinyl, and morpholinyl C<sub>2-8</sub> heterocyclic radical with at least one basic nitrogen atom in the ring, optionally including in the ring up to 3 additional heteroatoms or moieties independently selected from O, C=O, N, NH, NG, S, SO, and SO<sub>2</sub>, wherein G is R<sub>15</sub>, COR<sub>15</sub>, COOR<sub>15</sub>, SO<sub>2</sub>R<sub>15</sub>, SO<sub>2</sub>N, CSR<sub>15</sub>; or Z is NR<sub>11</sub>R<sub>12</sub> where each of R<sub>11</sub> and R<sub>12</sub> is independently selected from H, C<sub>1-6</sub> alkyl, phenyl, benzyl, C<sub>3-8</sub> cycloalkyl, and a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl, isoxazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indoliziny, isoindolyl, indolyl, indazolyl, purinyl, quinolyl, furazanyl, pyrrolidinyl, pyrrolinyl, imidazolidinyl, imidazoliny, pyrazolidinyl, pyrazolinyl, piperidyl, piperazinyl, indolinyl, and morpholinyl C<sub>2-5</sub> heterocyclic radical; or NR<sub>11</sub>R<sub>12</sub> taken together is a C<sub>6-8</sub> cycloalkylimino radical; and R<sub>15</sub> is C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkynyl, C<sub>2-6</sub> alkenyl, C<sub>3-7</sub> cycloalkyl, and C<sub>4-7</sub> cycloalkenyl; each of the above hydrocarbyl or heterocyclic groups being optionally substituted with between 1 and 3 substituents selected from C<sub>1-3</sub> alkyl, C<sub>1-3</sub> alkoxy, halo, hydroxy, phenyl, and phenyl(C<sub>1-3</sub> alkyl); and wherein each of the above heterocyclic groups may be

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attached to the rest of the molecule by a carbon atom or a heteroatom;

provided that  $R_b$ ,  $R_d$ ,  $R_5$ ,  $R_6$ ,  $R_7$  and  $R_8$ , if halo, are selected from chloro;

or a pharmaceutically acceptable salt, amide, ester, or hydrate thereof.

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2. (original) A compound of claim 1, wherein  $R_3$  is H or  $C_{1-3}$  alkyl.
3. (original) A compound of claim 1, wherein  $R_3$  is  $-(C=O)C_{1-6}$  alkyl.
4. (original) A compound of claim 1, wherein  $R_3$  is  $-SO_2(C_{1-3}$  alkyl).
5. (original) A compound of claim 4 wherein  $R_3$  is methylsulfonyl.
6. (original) A compound of claim 1, wherein  $W'$  is a covalent bond.
7. (original) A compound of claim 1, wherein  $W'$  is  $SO_2$  or  $(C=O)$ .
8. (original) A compound of claim 1, wherein  $R_c$  is WZ.
9. (original) A compound of claim 1, wherein  $R_b$  or  $R_d$  is WZ.
10. (original) A compound of claim 1, wherein W is ethoxy, propoxy, or butoxy.
11. (original) A compound of claim 1, wherein W is  $-O-$ .
12. (original) A compound of claim 1, wherein one of  $R_b$ ,  $R_c$ , and  $R_e$  is WZ and the others are independently selected from H, methyl, ethyl, methoxy, ethoxy, amino, nitro, and halo; and  $R_a$  and  $R_d$  are each independently H or methyl.

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13. (original) A compound of claim 1, wherein at least two of the following apply:  $R_c$  is WZ; W is propoxy or ethoxy; and Z is N-piperidino, 2-(N-methyl)pyrrolidino, or N,N-dimethyl.
14. (currently amended) A compound of claim 1, wherein Z is pyrrolidino, N-methyl-pyrrolidino, pyridyl, thiazoyl, piperidino, or  $NR_{11}R_{12}$  where each of  $R_{11}$  and  $R_{12}$  is independently selected from H, C<sub>1-6</sub> alkyl, phenyl, benzyl, C<sub>3-6</sub> cycloalkyl, and a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrol, imidazol, pyrazol, isothiazol, isoxazol, pyrid, pyrazin, pyrimidin, pyridazin, indolizin, isoindol, indol, indazol, purin, quinol, furazan, pyrrolidin, pyrrolin, imidazolidin, imidazolin, pyrazolidin, pyrazolin, piperid, piperazin, indolin, and morpholinC<sub>2-5</sub>-heterocyclic radical or taken together with the N form a C<sub>6-8</sub> cycloalkylamino radical.
15. (currently amended) A compound of claim 1, wherein one of  $R_b$ ,  $R_c$ , and  $R_e$  is WZ and the others are independently selected from H, methyl, ethyl, methoxy, ethoxy, amino, and halo; and  $R_a$  and  $R_d$  are each independently H or methyl;  
W is -O- or C<sub>1-3</sub> alkoxy;  
Z is pyrrolidino, N-methylpyrrolidino, pyridyl, thiazoyl, piperidino, piperazino, N-methylpiperazino, or  $NR_{11}R_{12}$  where each of  $R_{11}$  and  $R_{12}$  is independently selected from H, C<sub>1-2</sub> alkyl, phenyl, benzyl, C<sub>3-8</sub> cycloalkyl, and a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrol, imidazol, pyrazol, isothiazol, isoxazol, pyrid, pyrazin, pyrimidin, pyridazin, indolizin, isoindol, indol, indazol, purin, quinol, furazan, pyrrolidin, pyrrolin, imidazolidin, imidazolin, pyrazolidin, pyrazolin, piperid, piperazin, indolin, and morpholinC<sub>2-5</sub>-heterocyclic radical; each of  $R_6$  and  $R_7$  are each independently H, methyl, methoxy, or ethoxy; each of  $R_5$  and  $R_8$  is H.

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16. (original) A compound of claim 15, wherein  $R_3$  is H or  $-SO_2$  ( $C_{1-6}$  alkyl).
17. (original) A compound of claim 15, wherein  $R_3$  is  $SO_2$  (phenyl) and  $(C=O)(C_{1-6}$  alkyl).
18. (original) A compound of claim 15, selected from 2-[4-[2-[1-(methyl)-2-pyrrolidinyl]ethoxy]phenyl]-1H-indole, 2-[4-[2-[1-(methyl)-2-pyrrolidinyl]ethoxy]phenyl]-1-(methanesulfonyl)-1H-indole, and 2-[4-[3-piperidinopropoxy]phenyl]-1H-indole; ) 2-(4-(3-(4-methylpiperazinopropoxy)-phenyl)indole; and 1-(methanesulfonyl)-2-(4-(3-(4-methylpiperazinopropoxy)-phenyl)indole; or a pharmaceutically acceptable salt, amide, ester, or hydrate thereof.
19. (original) A compound of claim 15, selected from 2-[4-[3-piperidinopropoxy]phenyl]-1-(methanesulfonyl)-1H-indole, and 2-[3-[3-piperidinopropoxy]phenyl]-1-(methanesulfonyl)-1H-indole or a pharmaceutically acceptable salt, amide, ester, or hydrate thereof.
20. (original) A pharmaceutical composition comprising a compound of formula (I)B and a pharmaceutically acceptable carrier.
21. (currently amended) A pharmaceutical composition of claim 20, wherein said compound has a formula wherein: one of  $R_b$ ,  $R_c$ , and  $R_e$  is WZ and the others are independently selected from H, methyl, ethyl, methoxy, ethoxy, amino, and halo;  
 $R_a$  and  $R_d$  are each independently H or methyl;  
W is  $-O-$  or  $C_{1-3}$  alkoxy;  
Z is pyrrolidino, N-methylpyrrolidino, pyridyl, thiazoyl, piperidino, or  $NR_{11}R_{12}$  where each of  $R_{11}$  and  $R_{12}$  is independently selected from H,  $C_{1-2}$  alkyl, phenyl, benzyl,  $C_{3-8}$  cycloalkyl, and a heterocyclic radical selected from the group consisting of thiazoyl, furyl, pyran, isobenzofuranyl, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl,

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isoxazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indolizinyl,  
isoindolyl, indolyl, indazolyl, purinyl, quinolyl, furazanyl, pyrrolidinyl,  
pyrrolinyl, imdazolidinyl, imidazolynyl, pyrazolidinyl, pyrazolinyl,  
piperidyl, piperazinyl, indolinyl, and morpholinyl G-2-5-heterocyclic  
radical; and

R<sub>6</sub> and R<sub>7</sub> are each independently H, methyl, methoxy, or ethoxy.

22. (original) A pharmaceutical composition of claim 21, wherein said compound has a formula selected from 2-[4-[2-[1-(methyl)-2-pyrrolidinyl]ethoxy]phenyl]-1H-indole; 2-[4-[2-[1-(methyl)-2-pyrrolidinyl]ethoxy]phenyl]-1-(methylsulfonyl)-1H-indole; 2-[4-[3-Piperidinopropoxy]phenyl]-1H-indole; 2-[4-[3-Piperidinopropoxy]phenyl]-1-(methylsulfonyl)-1H-indole; 2-[3-[3-Piperidinopropoxy]phenyl]-1-(methylsulfonyl)-1H-indole; 2-(4-(3-(4-methylpiperazino)propoxy)-phenyl)indole; and 1-(methylsulfonyl)-2-(4-(3-(4-methylpiperazino)-propoxy)phenyl)indole; or a pharmaceutically acceptable salt, amide, ester, or hydrate thereof.
23. (cancelled)
24. (cancelled)
25. (cancelled)
26. (cancelled)
27. (cancelled)
28. (cancelled)
29. (cancelled)
30. (cancelled)